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Rachmady et al.

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(54) **SELECTIVE ETCH FOR PATTERNING A SEMICONDUCTOR FILM DEPOSITED NON-SELECTIVELY**

(58) **Field of Classification Search** 438/341, 438/349, 350, 489
See application file for complete search history.

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(*) Notice: Subject to any disclaimer, the term of this patent is extended or adjusted under 35 U.S.C. 154(b) by 0 days.

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This patent is subject to a terminal disclaimer.

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(22) Filed: **Feb. 20, 2008**

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(65) **Prior Publication Data**

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(57) **ABSTRACT**

Related U.S. Application Data

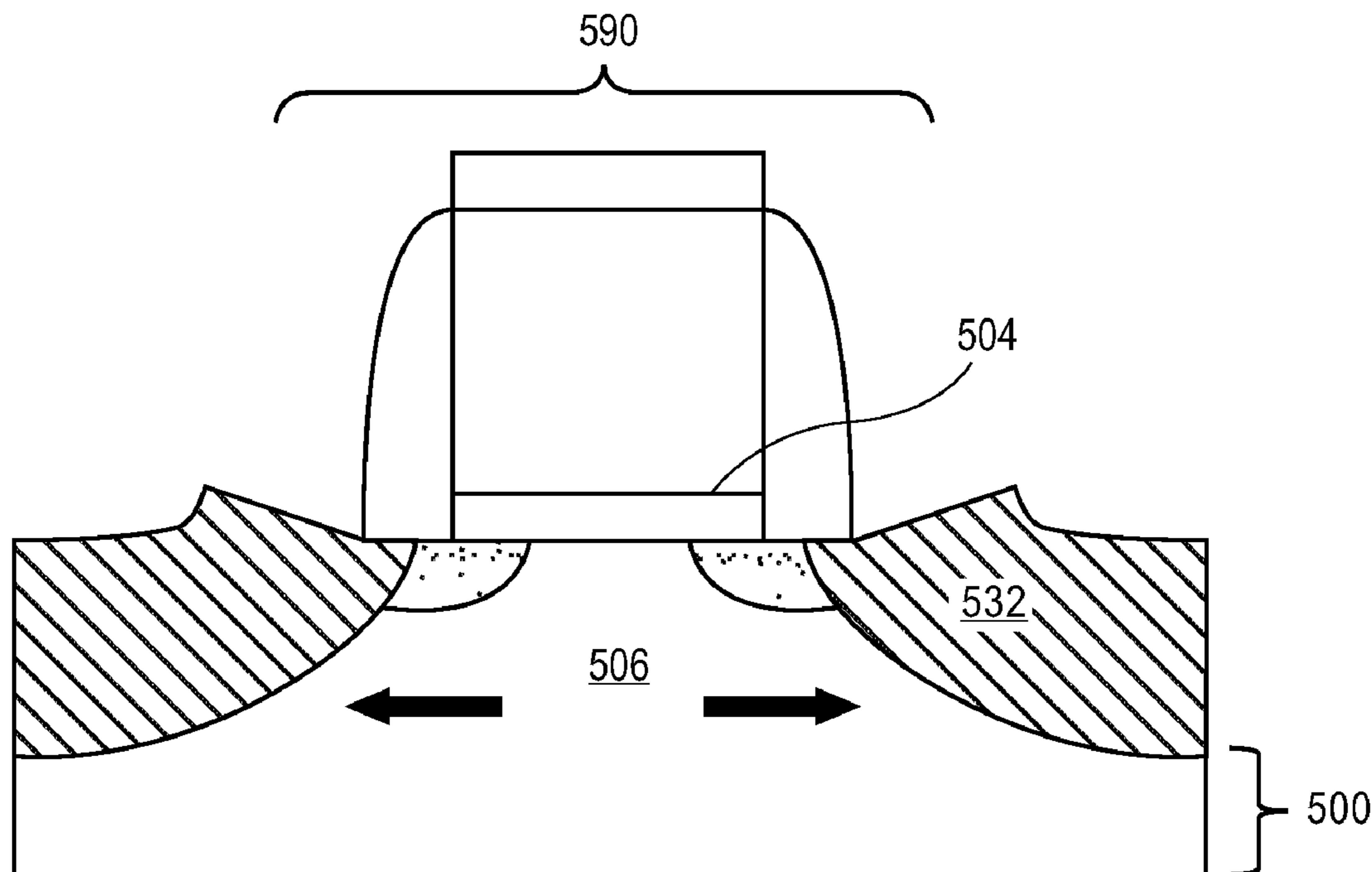
(63) Continuation of application No. 11/387,012, filed on Mar. 21, 2006, now Pat. No. 7,364,976.

A method to selectively etch, and hence pattern, a semiconductor film deposited non-selectively is described. In one embodiment, a carbon-doped silicon film is deposited non-selectively such that the film forms an epitaxial region where deposited on a crystalline surface and an amorphous region where deposited on an amorphous surface. A four-component wet etch mixture is tuned to selectively etch the amorphous region while retaining the epitaxial region, wherein the four-component wet etch mixture comprises an oxidizing agent, an etchant, a buffer and a diluent.

(51) **Int. Cl.**
H01L 21/20 (2006.01)

(52) **U.S. Cl.** **438/478**; 438/299; 438/300;
438/301; 438/482; 438/488; 257/E21.101;
257/E21.166

17 Claims, 9 Drawing Sheets



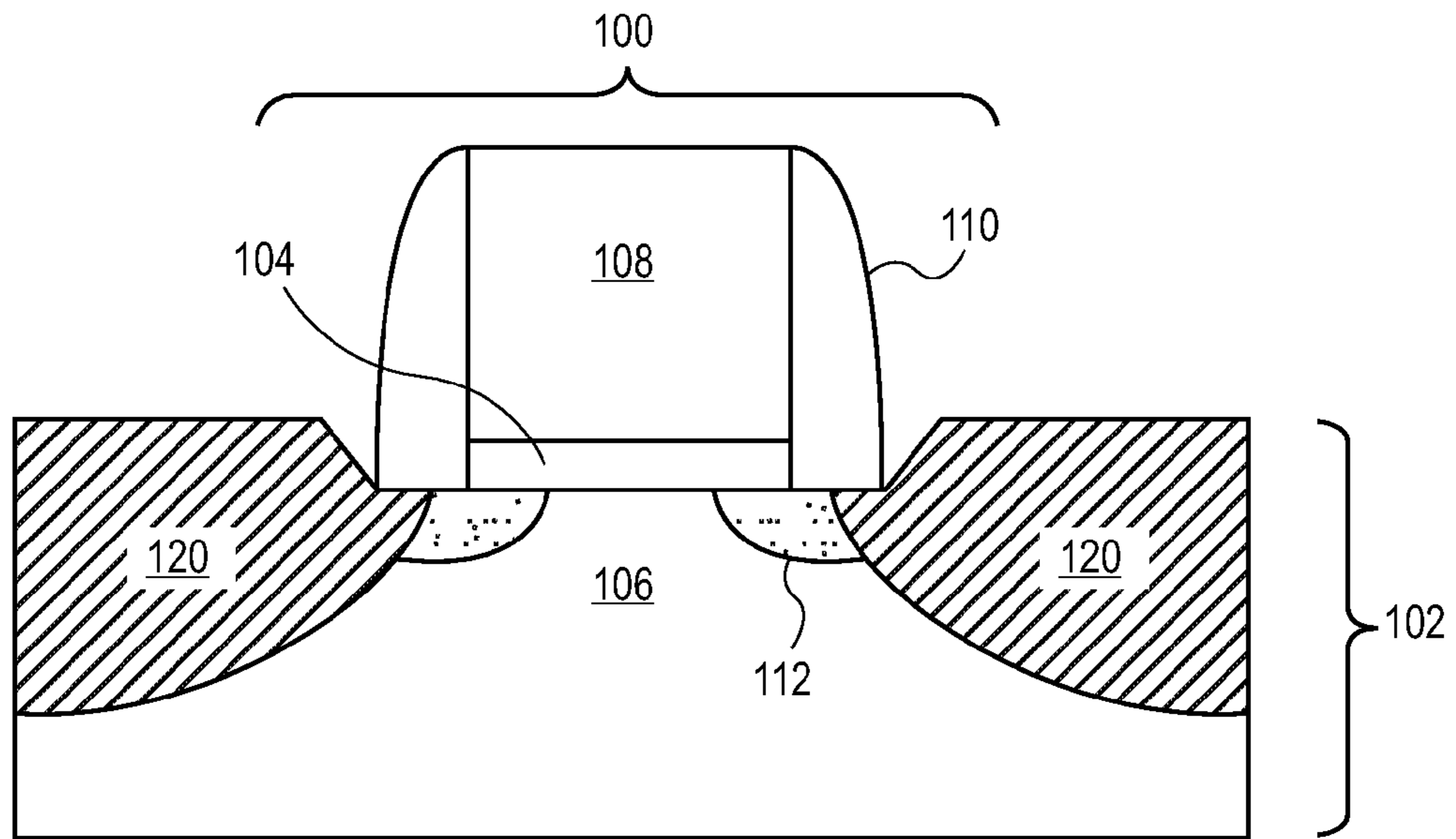


FIG. 1
(PRIOR ART)

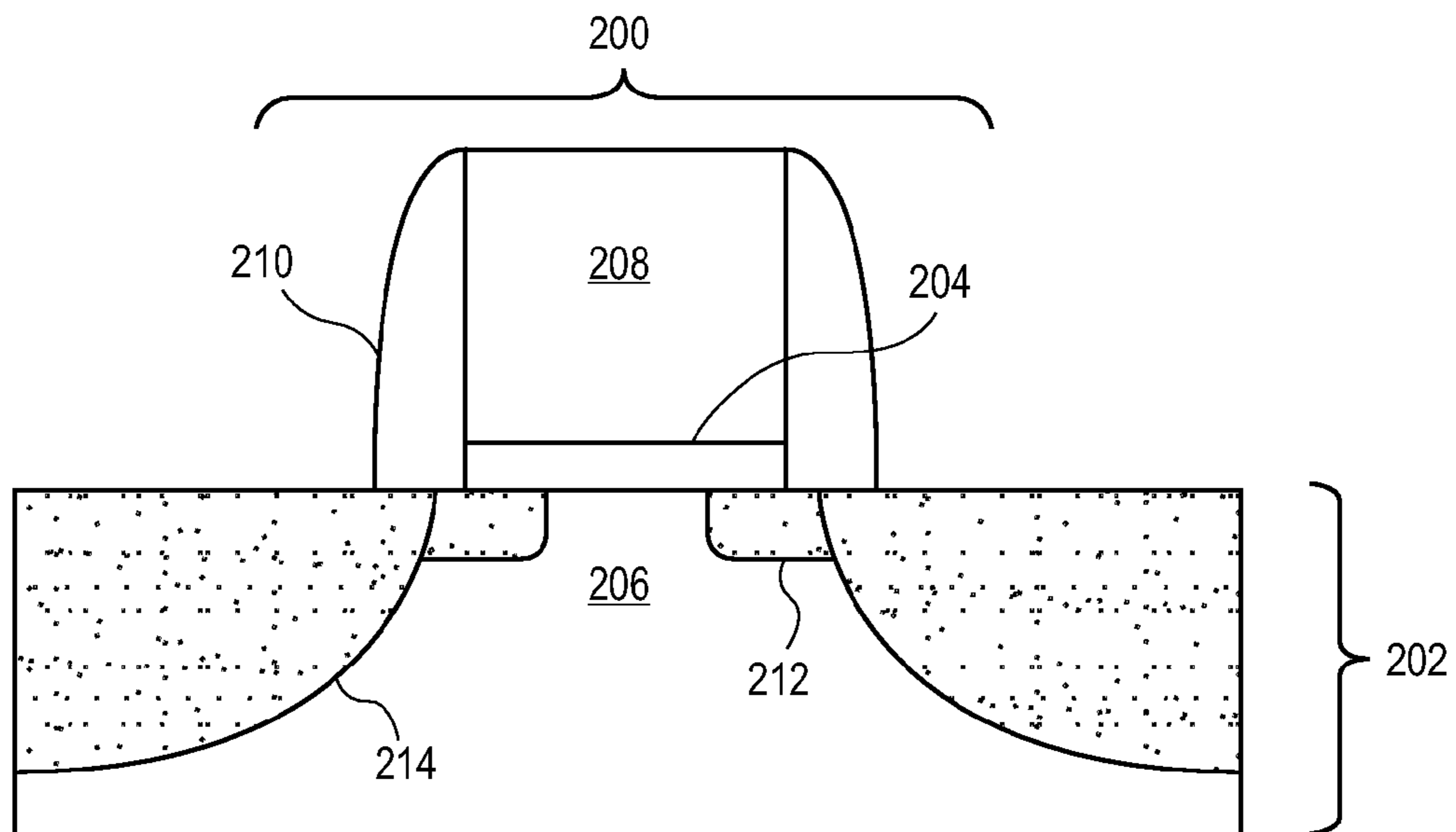


FIG. 2A
(PRIOR ART)

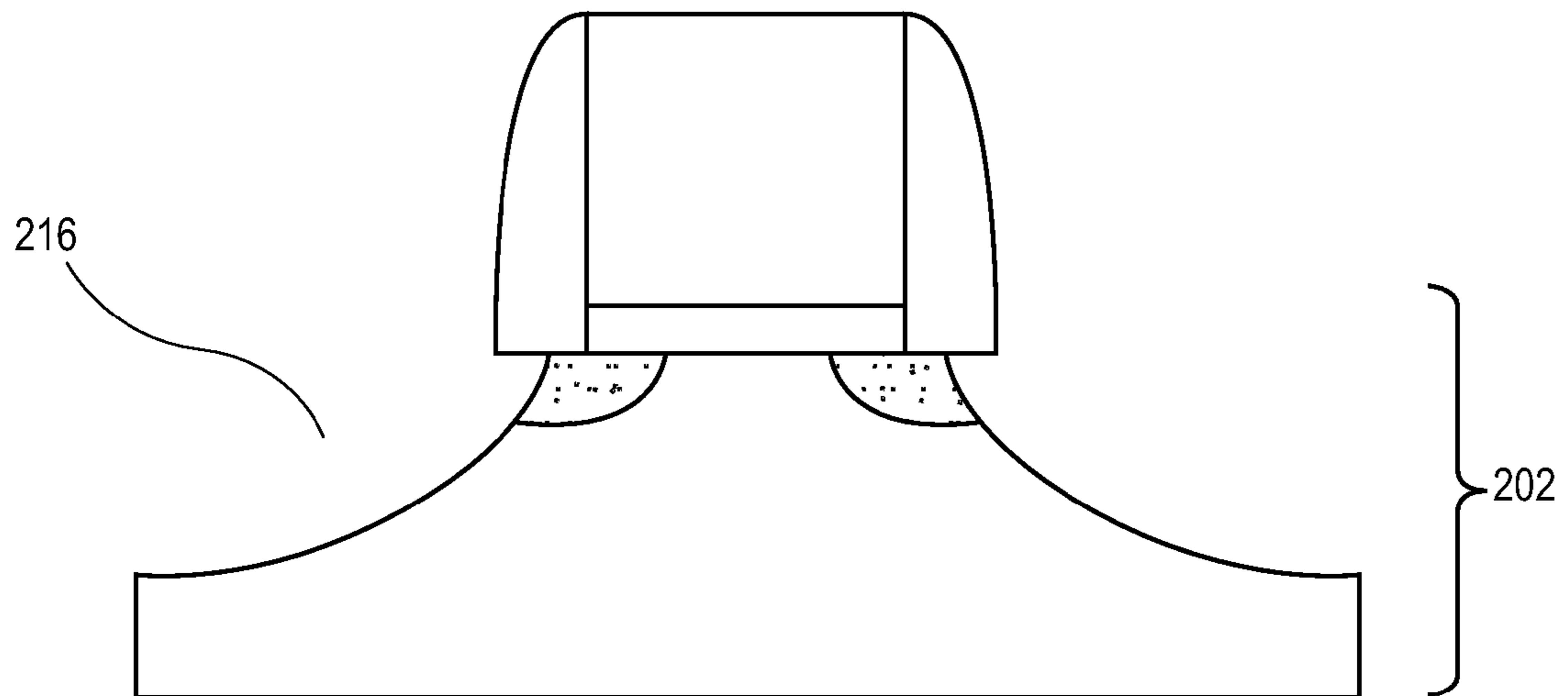


FIG. 2B

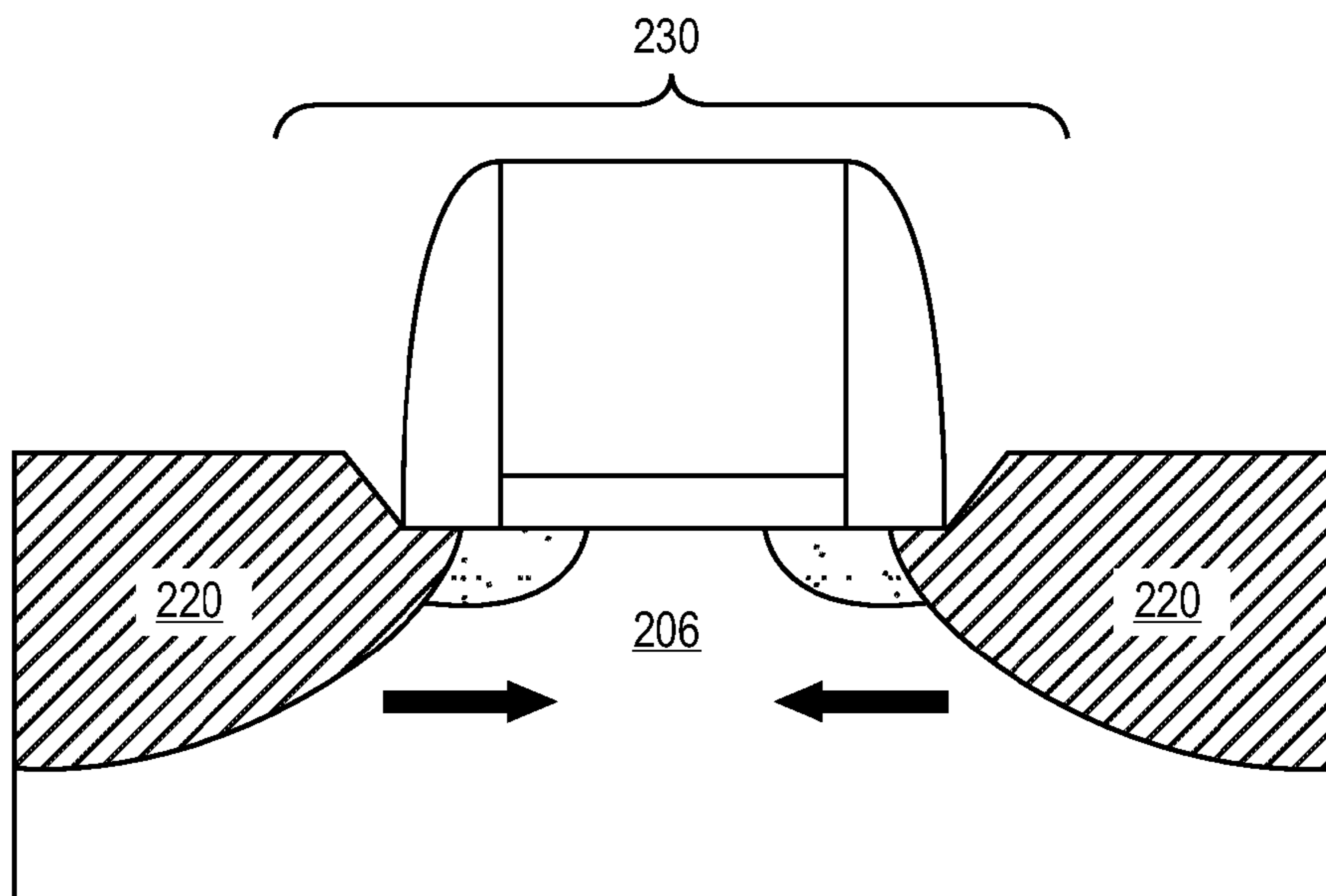


FIG. 2C

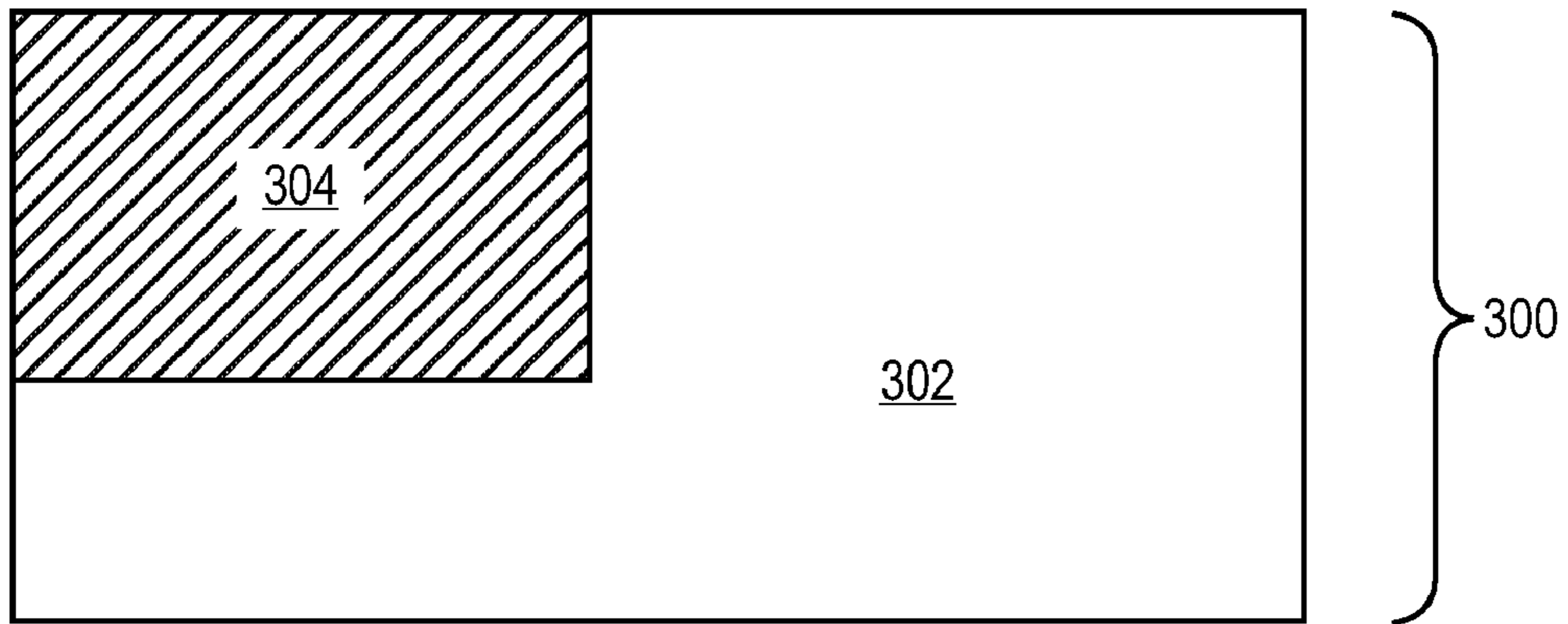


FIG. 3A

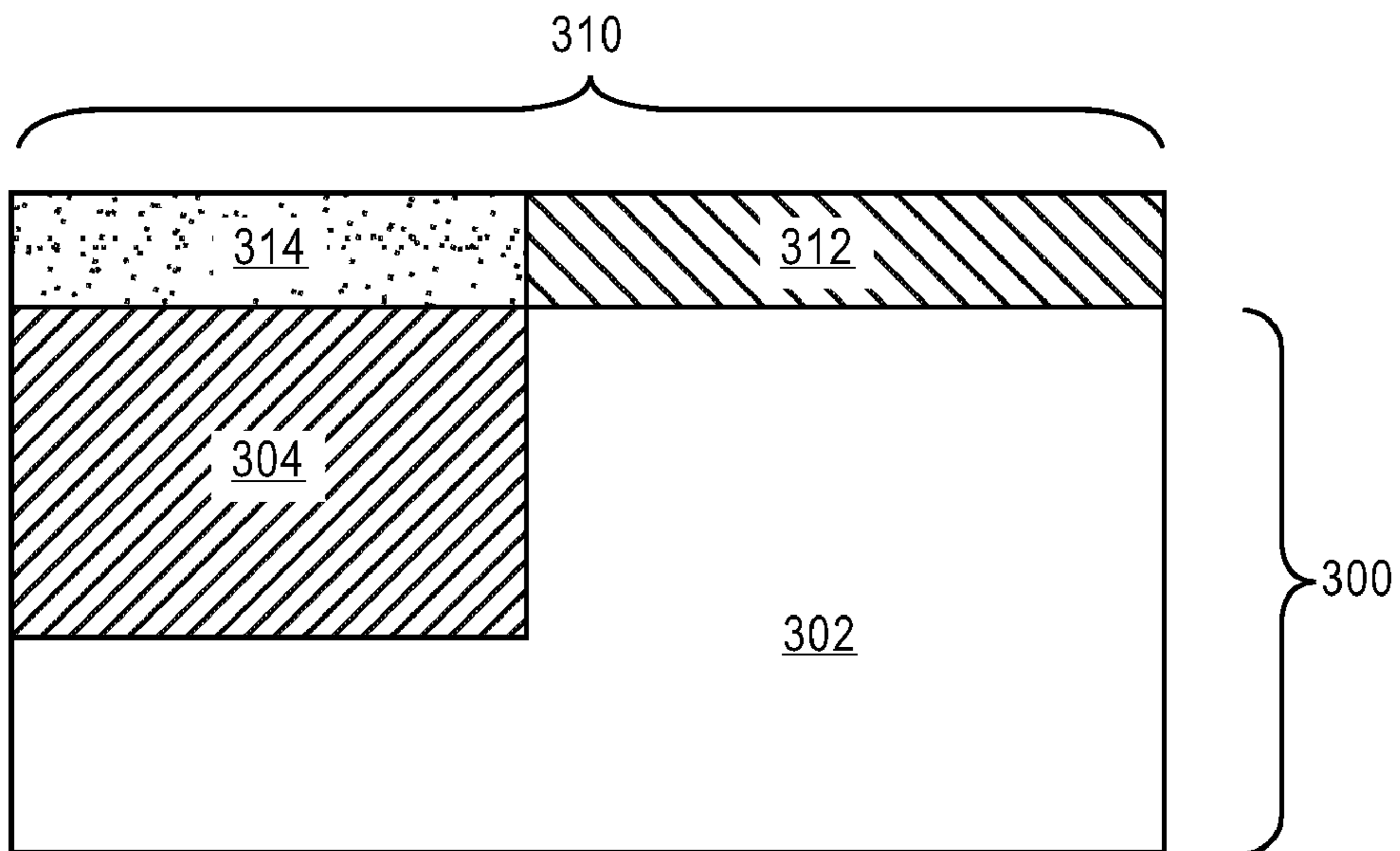


FIG. 3B

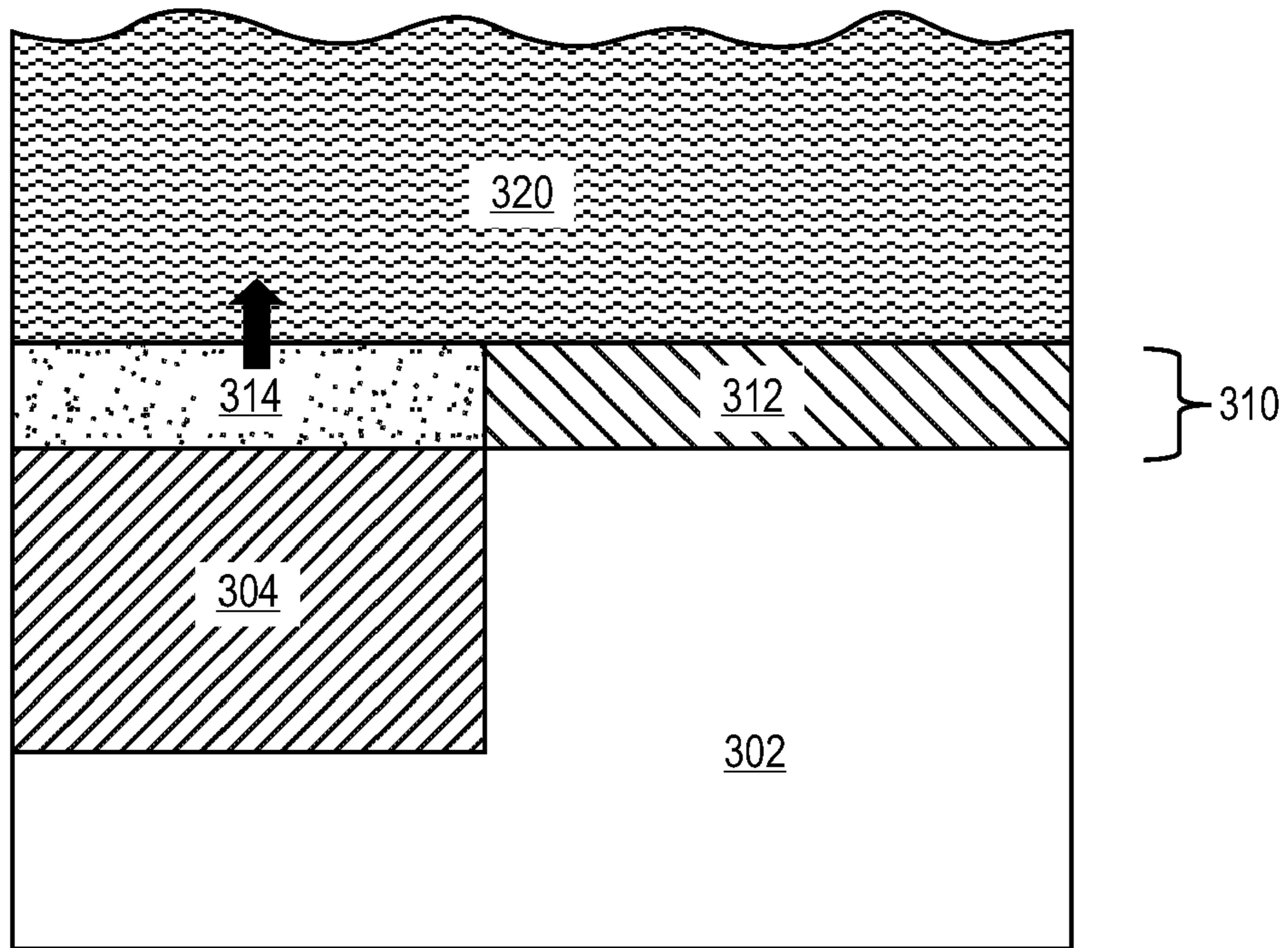


FIG. 3C

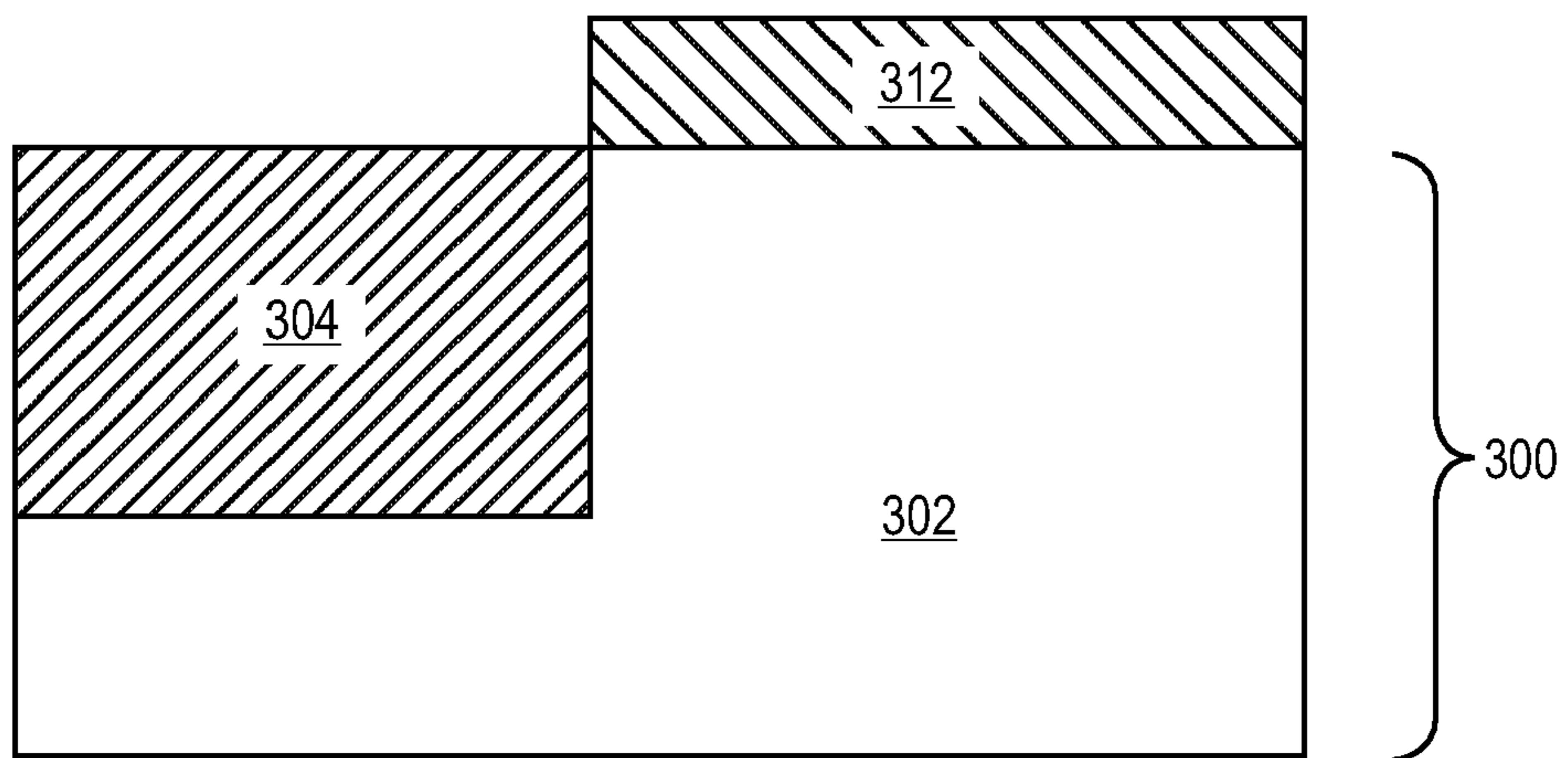


FIG. 3D

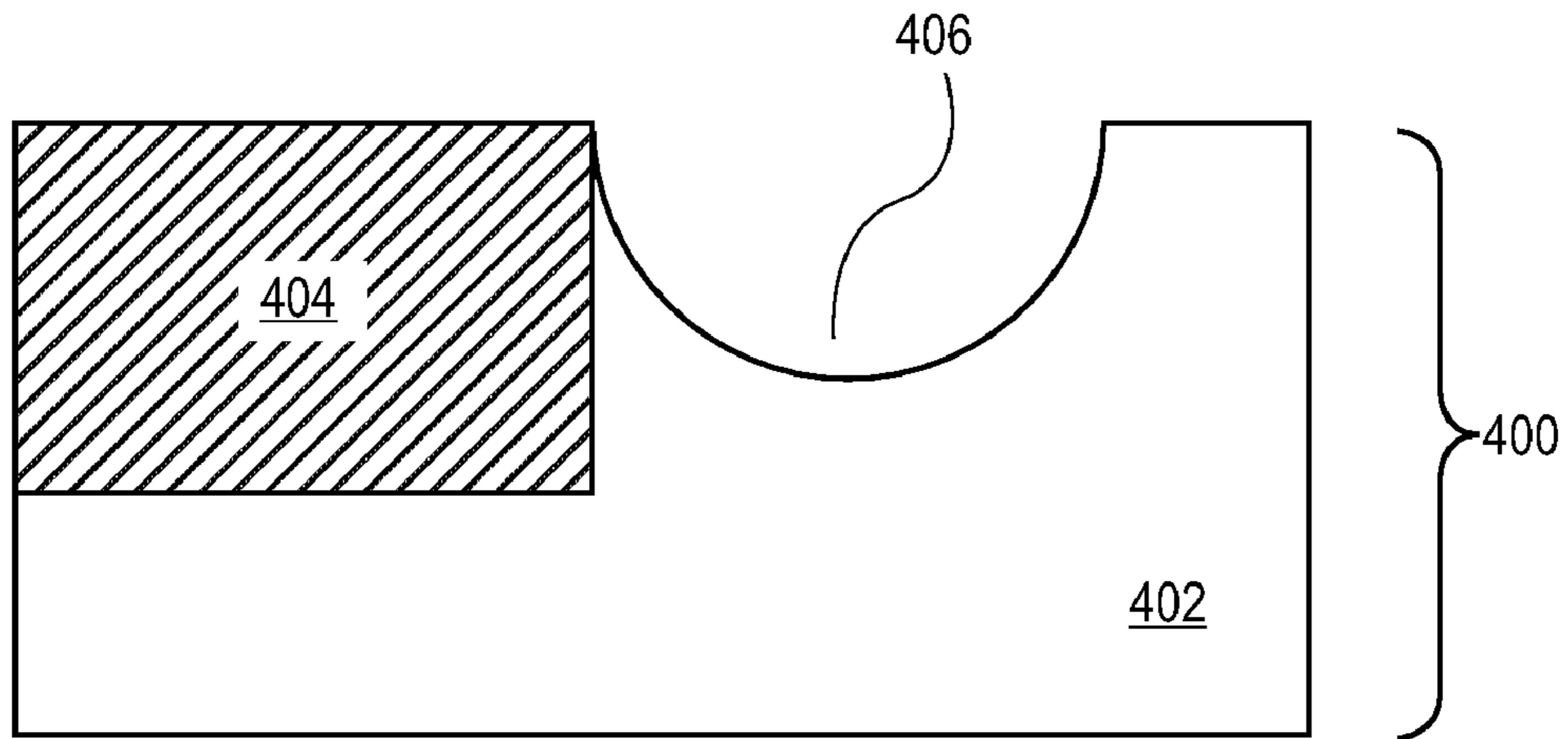


FIG. 4A

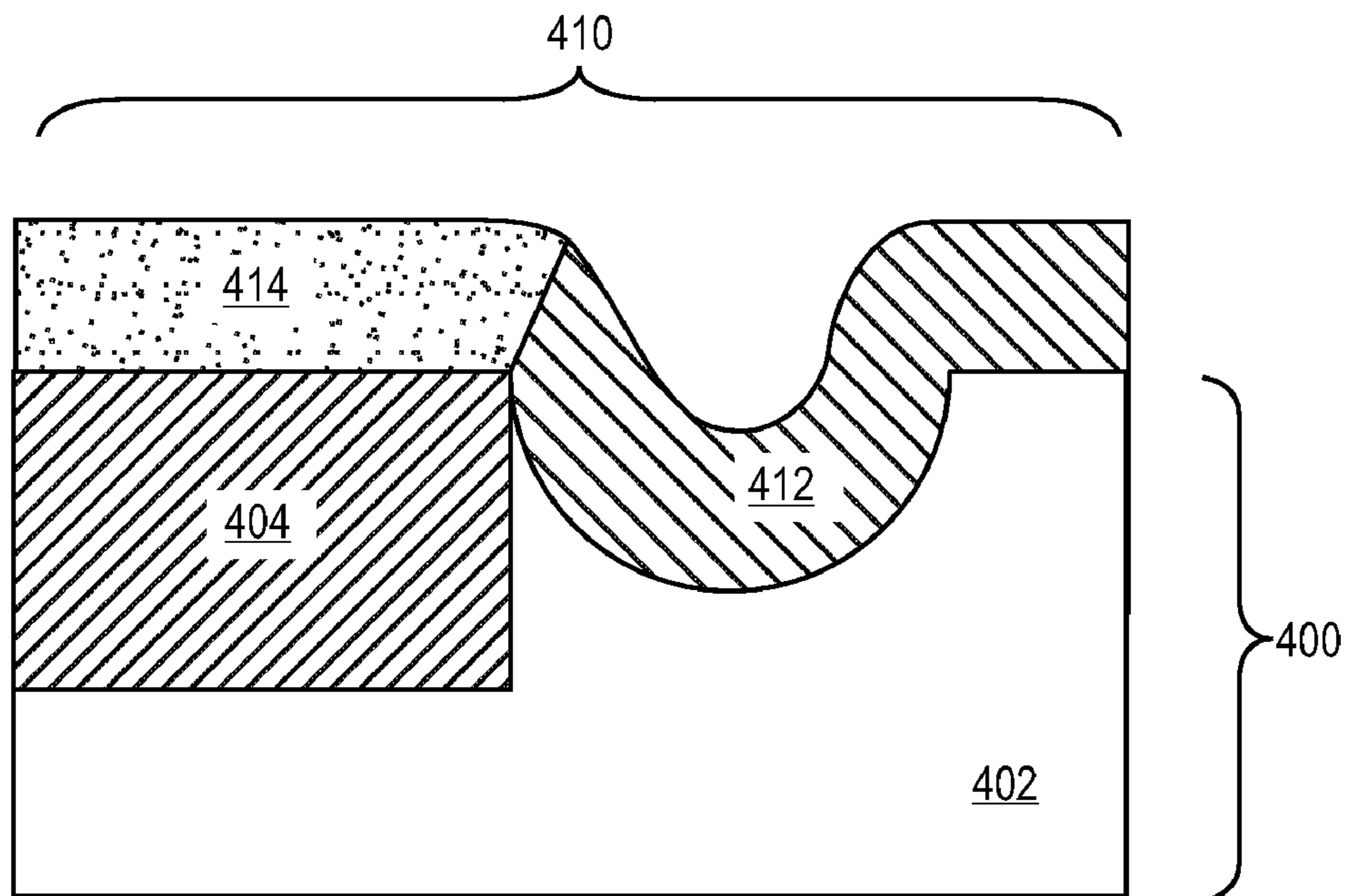


FIG. 4B

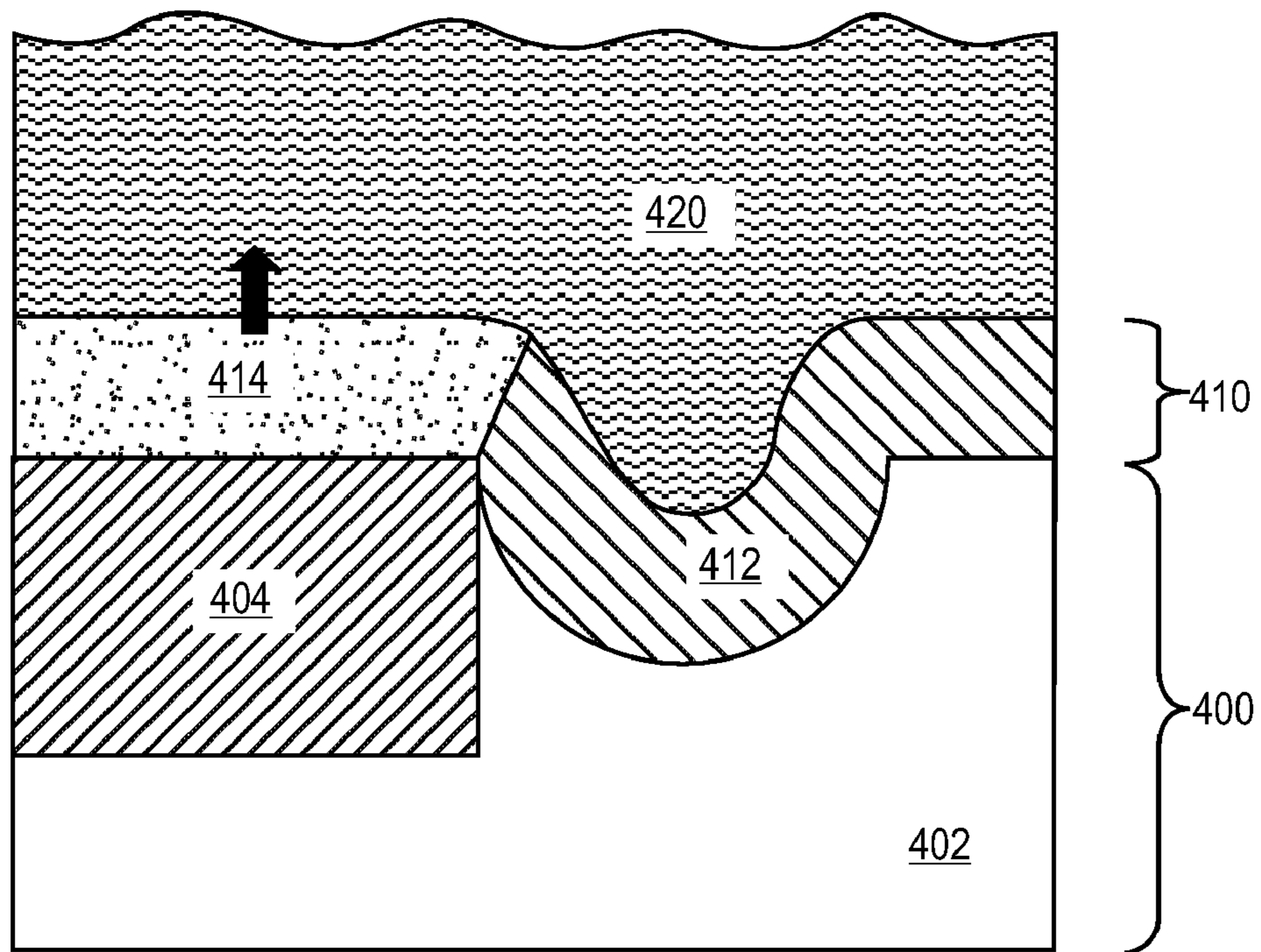


FIG. 4C

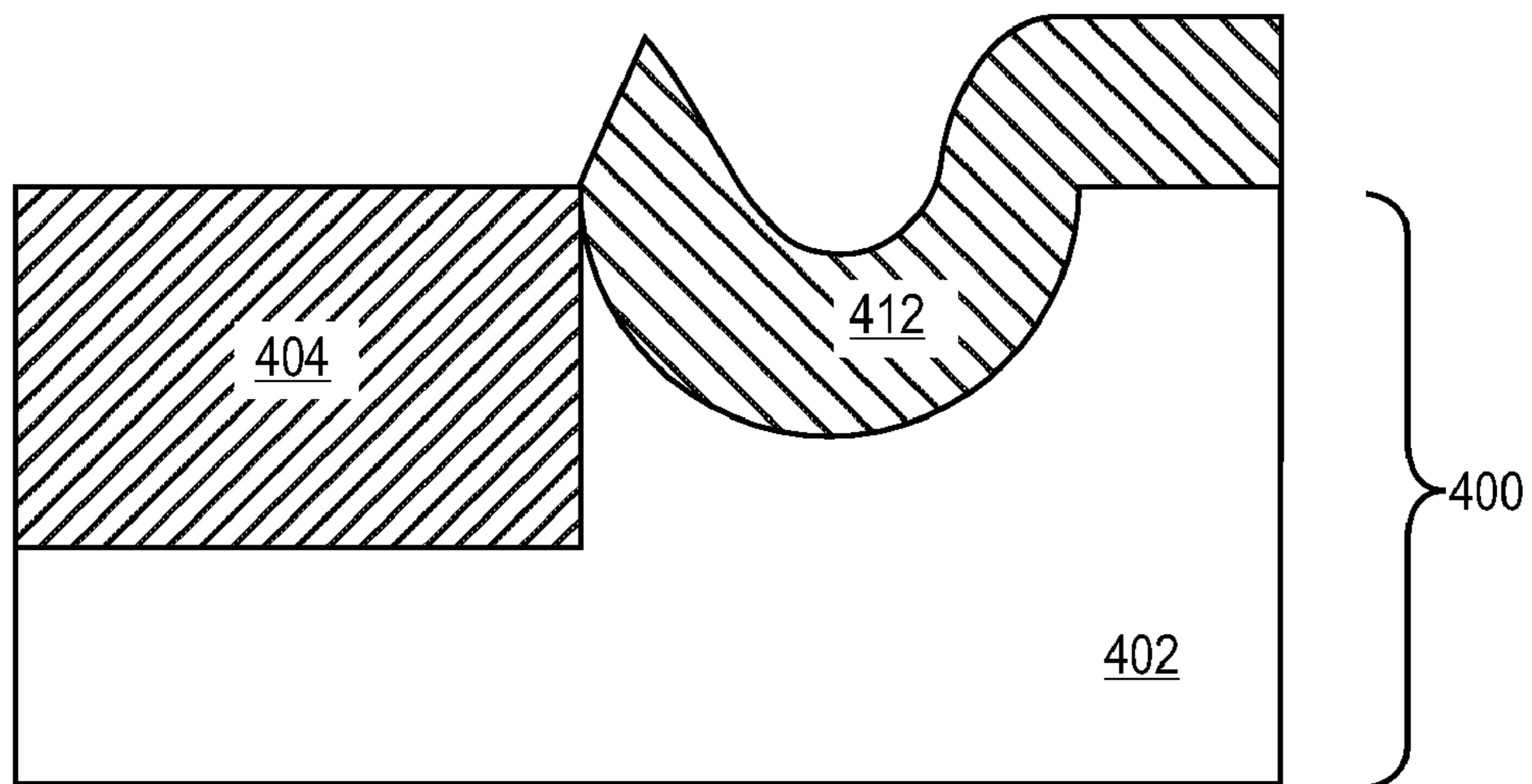


FIG. 4D

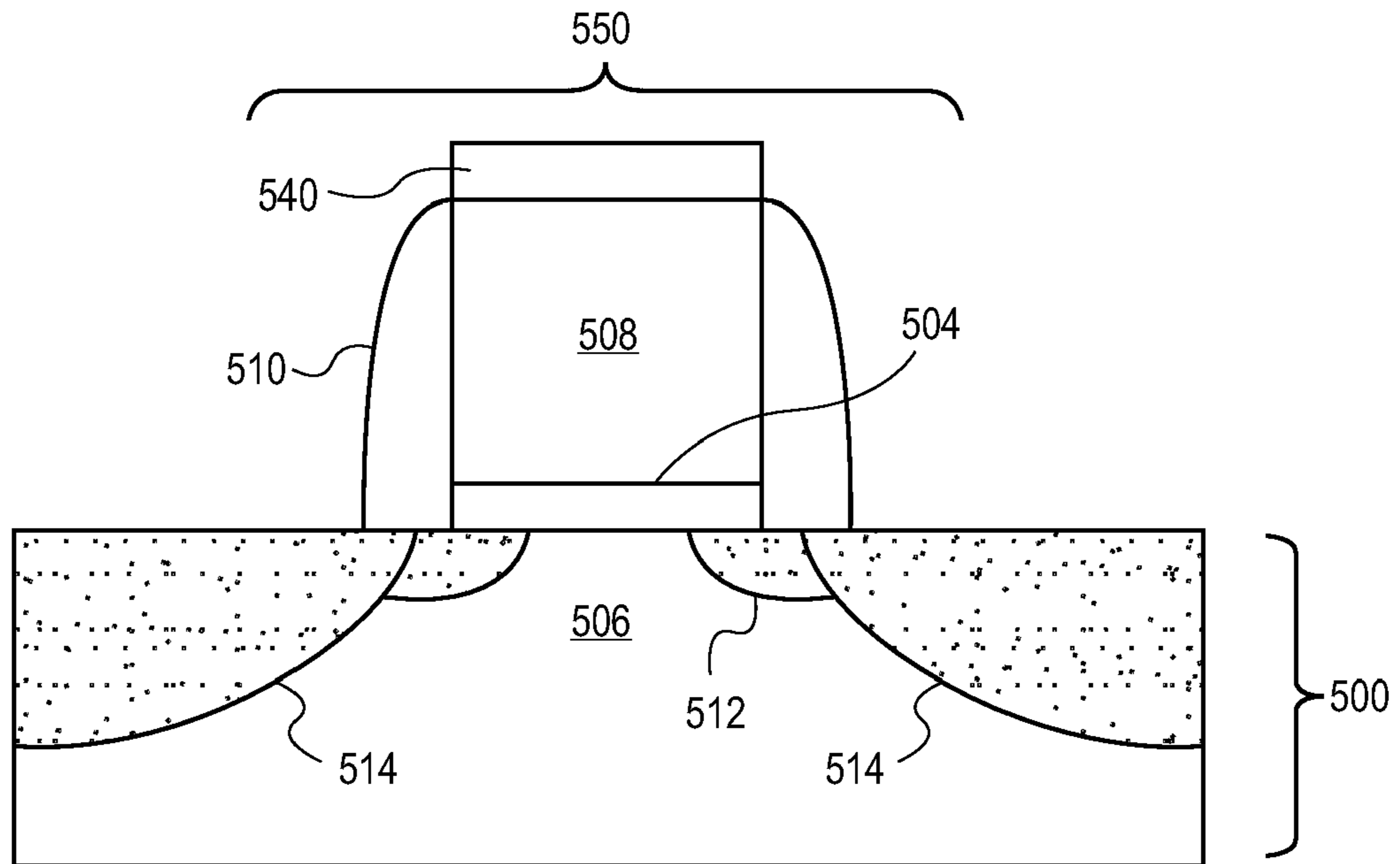


FIG. 5A

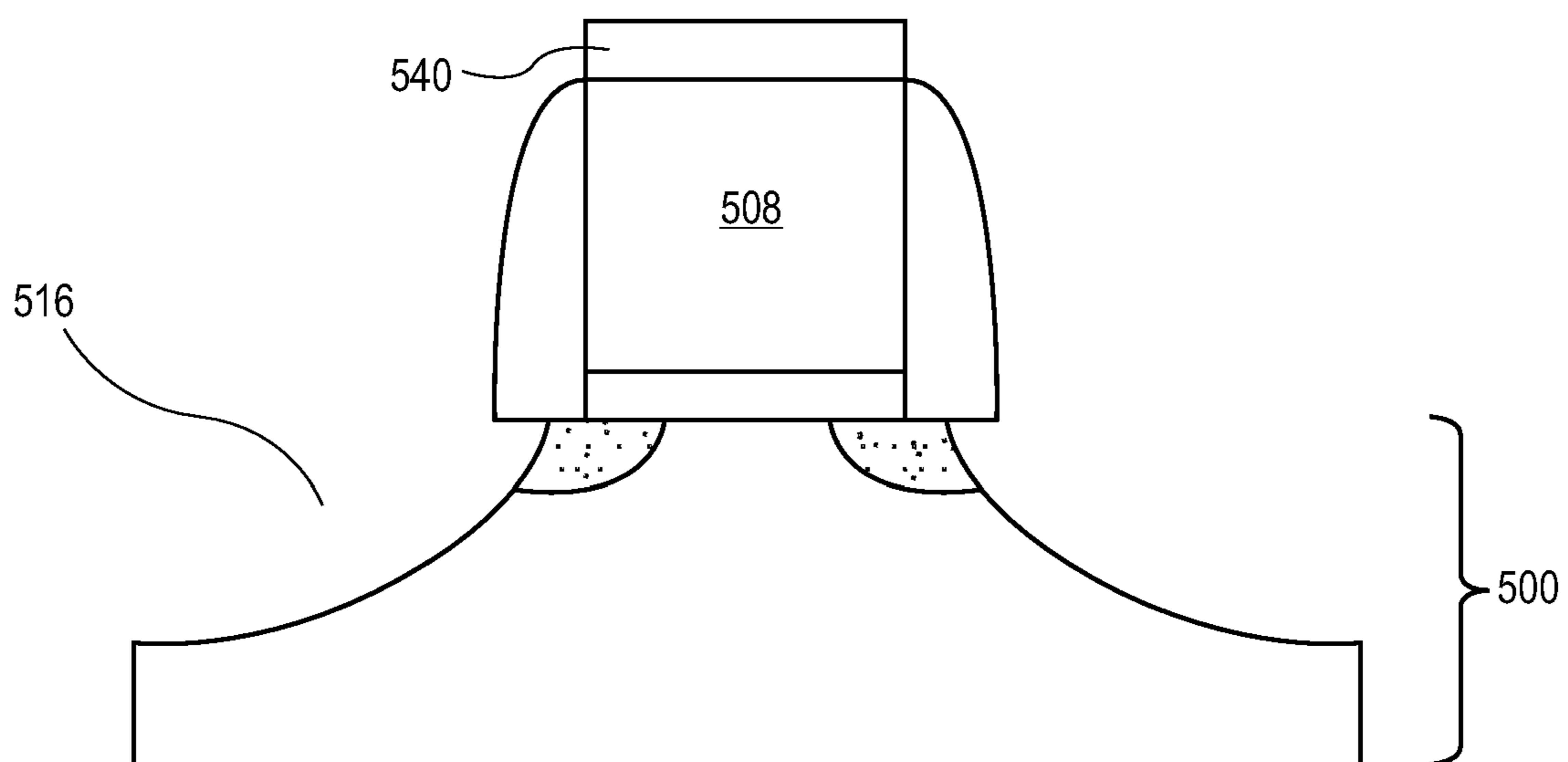


FIG. 5B

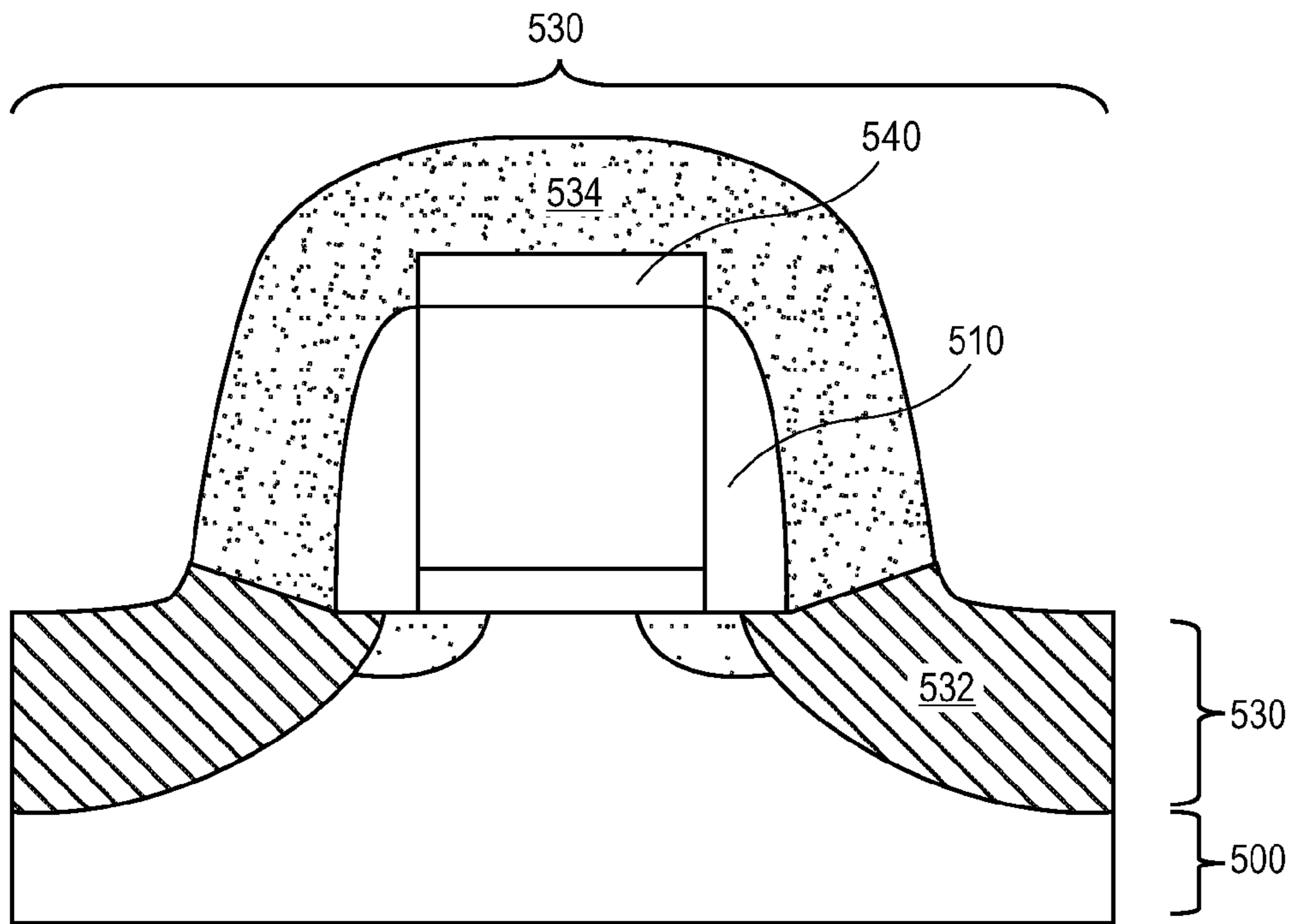


FIG. 5C

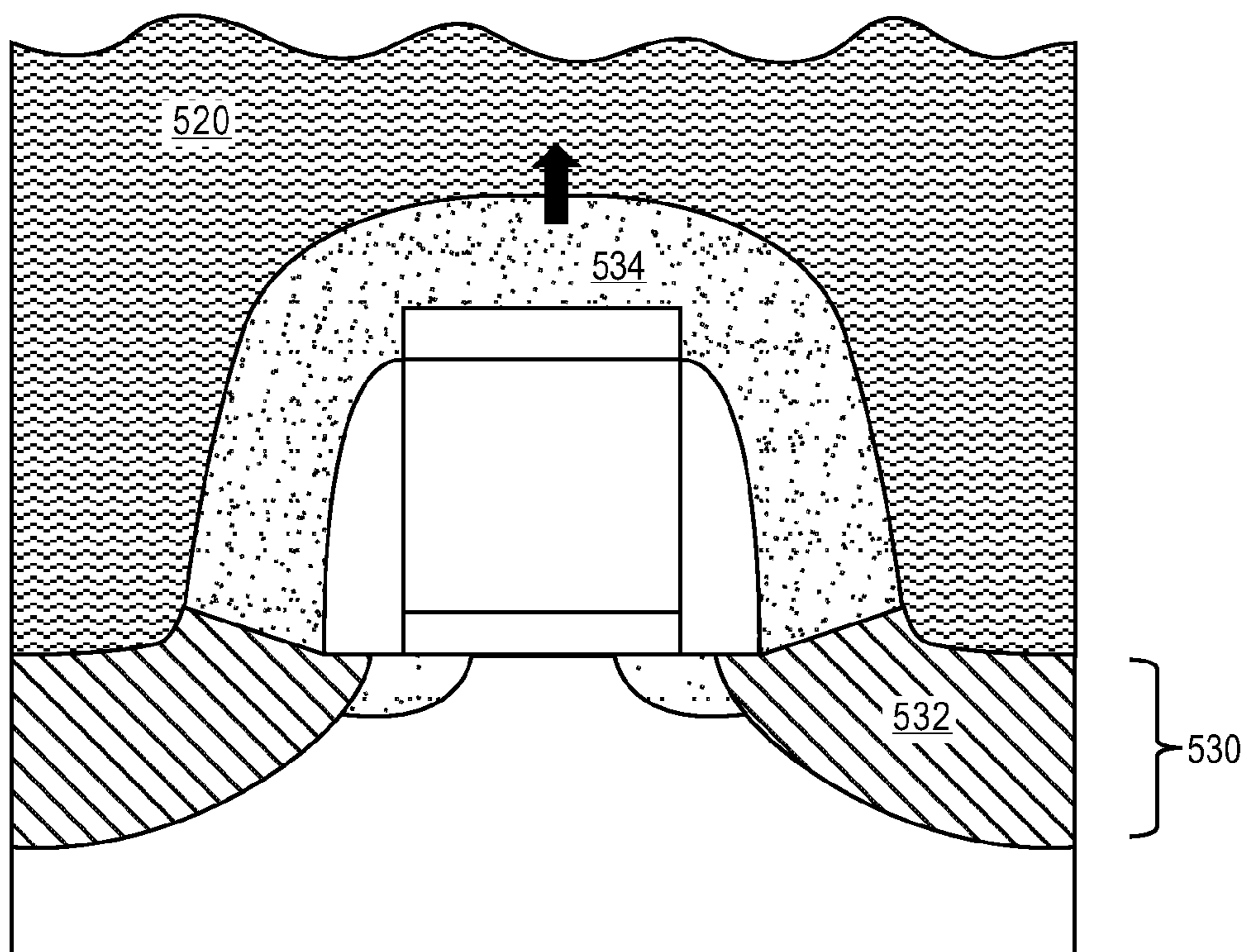


FIG. 5D

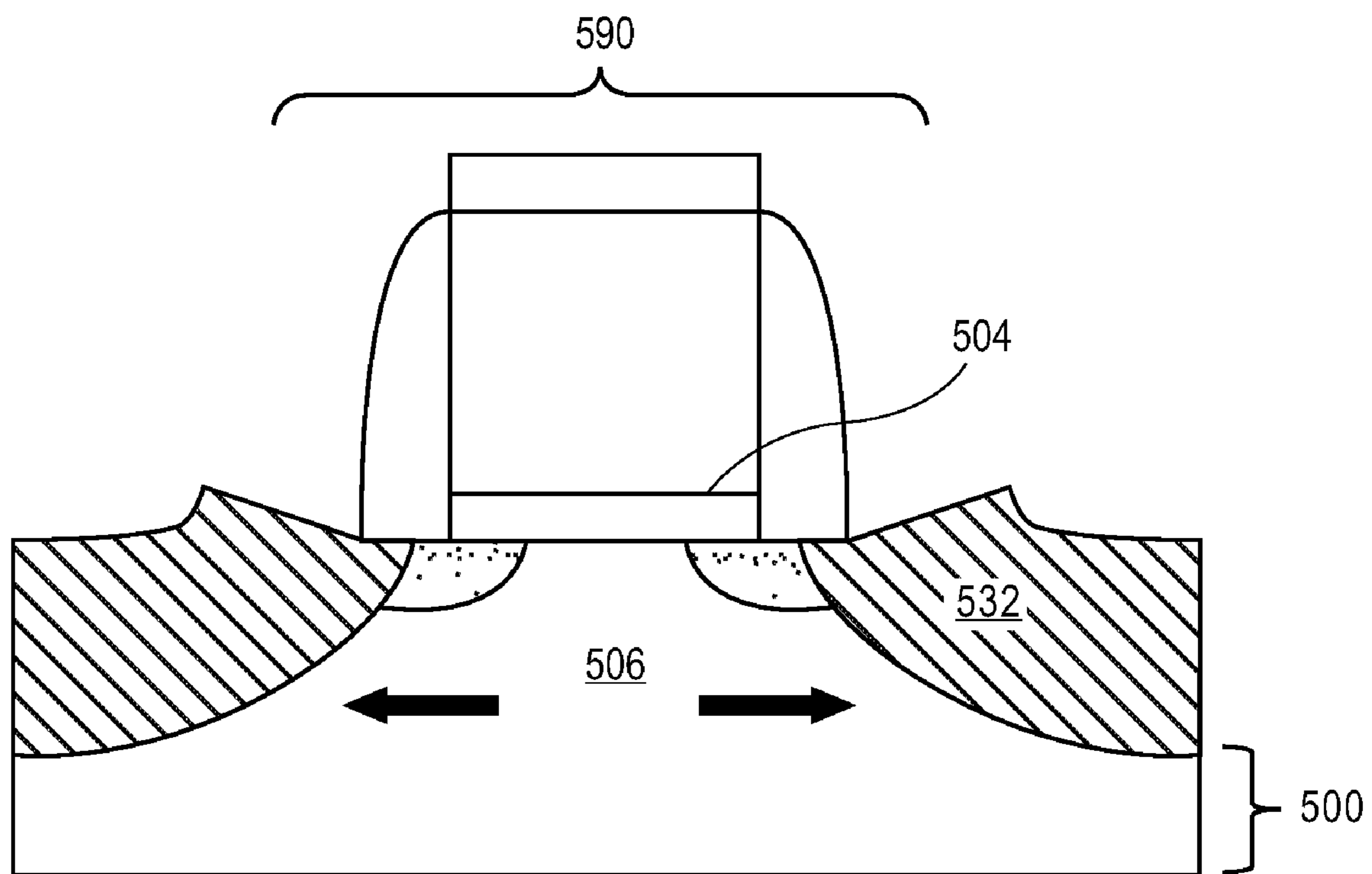


FIG. 5E

**SELECTIVE ETCH FOR PATTERNING A
SEMICONDUCTOR FILM DEPOSITED
NON-SELECTIVELY**

CROSS-REFERENCE TO RELATED
APPLICATIONS

This application is a continuation of U.S. patent application Ser. No. 11/387,012, filed Mar. 21, 2006, now U.S. Pat. No. 7,364,976 entitled "SELECTIVE ETCH FOR PATTERNING A SEMICONDUCTOR FILM DEPOSITED NON-SELECTIVELY," the entire contents of which are hereby incorporated by reference herein.

BACKGROUND OF THE INVENTION

1) Field of the Invention

The invention is in the field of Semiconductor Devices.

2) Description of Related Art

For the past several years, the performance of semiconductor devices, such as Metal Oxide Semiconductor Field-Effect Transistors (MOS-FETs), has been greatly enhanced by the incorporation of strained silicon regions into the active portions of a semiconductor substrate, e.g. the use of compressively strained silicon channel regions to enhance hole mobility in P-type Metal Oxide Semiconductor Field-Effect Transistors (PMOS-FETs). The presence of such strained silicon regions may greatly enhance the rate at which charge migrates in a channel when a semiconductor is in an ON state.

FIG. 1 depicts a typical strained PMOS-FET **100** fabricated on a substrate **102**. A gate dielectric layer **104** sits above a channel region **106** and a gate electrode **108** sits above a gate dielectric layer **104**. Gate dielectric layer **104** and gate electrode **108** are isolated by gate isolation spacers **110**. Tip extensions **112** are formed by implanting dopant atoms into substrate **102**. Strain-inducing source/drain regions **120** are formed by selectively growing an epitaxial film in etched-out portions of substrate **102** and are doped either in situ or after epitaxial film growth, or both. Strain-inducing source/drain regions are comprised of a material with a larger lattice constant than that of the channel region **106**. In typical PMOS-FETs, the channel region **106** is comprised of crystalline silicon, while the strain-inducing source/drain regions **120** are comprised of epitaxial silicon/germanium which has a larger lattice constant than that of crystalline silicon. Strain-inducing source/drain regions **120** can invoke a uniaxial compressive strain on the channel region **106**. Such a compressive strain in the channel region **106** can enhance the hole mobility in the channel region **106** of a PMOS-FET, lending to improved performance of the device.

FIGS. 2A-C illustrate a typical process flow for forming strain-inducing source/drain regions in a PMOS-FET. Referring to FIG. 2A, a non-strained PMOS-FET **200** is first formed. Non-strained PMOS-FET **200** is comprised of a channel region **206**. A gate dielectric layer **204** sits above the channel region **206** and a gate electrode **208** sits above gate dielectric layer **204**. Gate dielectric layer **204** and gate electrode **208** are isolated by gate isolation spacer **210**. Tip extensions **212** and source/drain regions **214** are formed by implanting dopant atoms into substrate **202**. Thus, the source/drain regions **214** are initially formed from the same material as the channel region **206**. Therefore, the lattice mismatch between the source/drain regions **214** and the channel region **206** is negligible, resulting in effectively no strain on the channel region **206**. Referring to FIG. 2B, portions of substrate **202** are removed, e.g. by an etch process, to form recessed regions **216** in substrate **202**. Subsequently, strain-

inducing source/drain regions **220** are formed by selectively growing an epitaxial film into recessed regions **216**, as depicted in FIG. 2C. Strain-inducing source/drain regions **220** can be doped with charge carrier atoms, e.g. boron in the case of a PMOS-FET, which may be done in situ (during the deposition of the epitaxial film) or after epitaxial film growth (which may require a subsequent anneal process), or both. In an example, substrate **202**, and hence channel region **206**, is comprised of crystalline silicon and the film grown to form strain-inducing source/drain regions **220** is comprised of epitaxial silicon/germanium. The lattice constant of the epitaxial silicon/germanium film can be greater than that of crystalline silicon by a factor of ~1% (for 70% Si, 30% Se) and so strain-inducing source/drain regions **220** are comprised of a material with a larger lattice constant than that of the channel region **206**. Therefore, a uniaxial compressive strain, depicted by the arrows in FIG. 2C, is rendered on channel region **206** in PMOS-FET **230**, which can enhance hole mobility in the device.

In order to improve performance in N-type Metal Oxide Semiconductor Field-Effect Transistors (NMOS-FETs), a uniaxial tensile strain may be required to enhance electron mobility in the channel region. This may require incorporation of strain-inducing source/drain regions with a smaller lattice constant than that of the channel region. For example, epitaxial carbon-doped silicon source/drain regions may be desirable for NMOS-FETs with a crystalline silicon channel region because the lattice constant of epitaxial carbon-doped silicon is smaller than that of crystalline silicon. However, selective deposition of an epitaxial carbon-doped silicon film can be difficult to control with very narrow processing windows. Thus, a method to pattern a non-selective N-type epitaxial carbon-doped silicon film is described herein.

BRIEF DESCRIPTION OF THE DRAWINGS

FIG. 1 illustrates a cross-sectional view of a strained P-type Metal Oxide Semiconductor Field-Effect Transistor (PMOS-FET), in accordance with the prior art.

FIGS. 2A-C illustrate cross-sectional views representing the formation of a PMOS-FET with strain-inducing source/drain regions, in accordance with the prior art.

FIGS. 3A-D illustrate cross-sectional views representing the non-selective deposition and subsequent selective etching of a semiconductor film, in accordance with an embodiment of the present invention.

FIGS. 4A-D illustrate cross-sectional views representing the non-selective deposition and subsequent selective etching of a semiconductor film on a substrate comprising a crystalline portion and an amorphous portion, wherein a region in the crystalline portion is recessed to form an etched-out region in the crystalline portion, in accordance with an embodiment of the present invention.

FIGS. 5A-E illustrate cross-sectional views representing a process flow for forming an NMOS-FET device with strain-inducing source/drain regions as facilitated by a four-component selective etch mixture, in accordance with an embodiment of the present invention.

DETAILED DESCRIPTION

A process for fabricating semiconductor devices and the resultant devices are described. In the following description, numerous specific details are set forth, such as specific dimensions and chemical regimes, in order to provide a thorough understanding of the present invention. It will be apparent to one skilled in the art that the present invention may be

practiced without these specific details. In other instances, well-known processing steps, such as patterning steps, are not described in detail in order to not unnecessarily obscure the present invention. Furthermore, it is understood that the various embodiments shown in the figures are illustrative representations and are not necessarily drawn to scale.

Disclosed herein is a method to selectively etch, and hence pattern, a semiconductor film deposited non-selectively. Selective etching to pattern a semiconductor film after its non-selective deposition may enable the use of a broader process window at the time of deposition because the constraints associated with selective deposition are removed. For example, in accordance with an embodiment of the present invention, a carbon-doped silicon film is deposited non-selectively such that the film forms an epitaxial region where deposited on a crystalline surface and an amorphous region where deposited on an amorphous surface. By not being confined to a selective deposition such that deposition occurs only on crystalline surfaces, a greater variety of carbon-doped silicon films may be utilized, e.g. a broader range of films with varying carbon composition may be accessible. However, it may be desirable to retain only the crystalline (epitaxial) regions of a non-selectively deposited semiconductor film.

A selective etch process may selectively remove the amorphous regions of a non-selectively deposited semiconductor film, while retaining the crystalline regions of the non-selectively deposited semiconductor film. Depending on the composition of the specific film being selectively etched, such a selective etch process may require tuning to accommodate the characteristics of a specific film. For an optimized etch process, a selectivity of greater than 10:1, and even greater than 20:1, may be achieved when etching an amorphous component of a film while retaining an epitaxial portion of a film.

A four-component wet etch mixture may be amenable to the necessary tunability required to accommodate the selective etching of a variety of film compositions, wherein the four-component wet etch mixture comprises an oxidizing agent, an etchant, a buffer and a diluent. An oxidizing agent may be employed to modify a portion of a film for selective etching by converting that portion to its corresponding oxide. For example, an oxidizing agent may oxidize a portion of a silicon film to silicon dioxide. In accordance with an embodiment of the present invention, the oxidizing agent is nitric acid, hydrogen peroxide or di-tert-butylperoxide. An etchant may be employed to dissolve the portion of the film that was oxidized by the oxidizing agent. For example, an etchant may dissolve only the portion of a silicon film that was oxidized by the oxidizing agent to form silicon dioxide. In accordance with an embodiment of the present invention, the etchant is hydrofluoric acid, ammonium fluoride, or tetramethyl ammonium fluoride. A buffer may be used to mitigate potential pH swings that may otherwise occur as component of the etched film become dissolved in the four-component wet etch mixture, enabling maintenance of a set pH within a factor of 0.1. In accordance with an embodiment of the present invention, the buffer is acetic acid, methanol, or ethanol. A diluent may be utilized to slow the activity, and hence etch rate, of a four-component wet etch mixture, enabling control over the timing of the etch process. In accordance with an embodiment of the present invention, the diluent is water and it is either added to the four-component wet etch mixture as a distinct ingredient or is already present as a component of the oxidizing agent, the etchant, or the buffer. In one embodiment, the four component wet etch mixture exhibits etch selectivity greater than 20:1 between amorphous and crystalline portions

of a film, meaning that the amorphous portion is removed at least 20-fold faster than the crystalline portion.

For example, in accordance with one embodiment of the present invention, the semiconductor film to be selectively etched (i.e. amorphous regions removed and epitaxial regions retained) comprises a carbon-doped silicon film and the four-component wet etch mixture comprises an oxidizing agent (nitric acid), an etchant (hydrofluoric acid), a buffer (acetic acid), and a diluent (water). As the % carbon composition varies in non-selectively deposited carbon-doped silicon film, the local pH (the pH of the first several mono-layers of the wet etch mixture adjacent the carbon-doped silicon film) may vary, potentially compromising the selectivity of the wet etch mixture. Thus, in order to accommodate the characteristics of a specific non-selectively deposited film during a selective etch process, the component ratios of the four-component wet etch mixture may require modification to maintain etch selectivity between the amorphous regions and the epitaxial regions of the film.

As an example of one embodiment of the present invention, FIGS. 3A-D illustrate the non-selective deposition and subsequent selective etching of a semiconductor film. Referring to FIG. 3A, a suitable substrate **300** may comprise a single crystalline portion **302** and an amorphous portion **304**. In one embodiment of the present invention, crystalline portion **302** is comprised of crystalline silicon or an epitaxial silicon region grown atop a distinct crystalline silicon substrate. In another embodiment, crystalline portion **302** is comprised of germanium or a III-V material such as but not limited to gallium nitride, gallium phosphide, gallium arsenide, indium phosphide or indium antimonide. In one embodiment, amorphous portion **304** is comprised of an oxide or a nitride material. In another embodiment, amorphous portion **304** is comprised of silicon nitride, silicon dioxide, silicon oxy-nitride or a high-k material such as hafnium oxide.

Referring to FIG. 3B, semiconductor film **310** may be deposited non-selectively on the crystalline portion **302** and on the amorphous portion **304** of substrate **300**. In accordance with one embodiment of the present invention, a suitable non-selectively deposited semiconductor film **310** is one which forms an epitaxial region **312** above crystalline portion **302** and an amorphous region **314** above amorphous portion **304**. Semiconductor film **310** may be deposited by any suitable technique. In one embodiment, semiconductor film **310** is deposited by chemical vapor deposition, physical vapor deposition or atomic layer deposition. In one embodiment, semiconductor film **310** is lattice-matched with crystalline portion **302**. In another embodiment, semiconductor film **310** is lattice-mismatched with and has a smaller lattice constant than crystalline portion **302**. In another embodiment, semiconductor film **310** is lattice-mismatched with and has a larger lattice constant than crystalline portion **302**. In an embodiment, semiconductor film **310** is a carbon-doped silicon film. In another embodiment, semiconductor film **310** is a carbon-doped silicon film in situ doped with phosphorus dopant atoms. In the case of a carbon-doped silicon film, the carbon concentration as a % of total film composition may vary depending on the required application. In one embodiment, the total carbon concentration of a carbon-doped silicon semiconductor film is 0-0.1%. In another embodiment, the total carbon concentration of a carbon-doped silicon semiconductor film is 0.1-2%. In another embodiment, the total carbon concentration of a carbon-doped silicon semiconductor film is greater than 2%. In one embodiment, a carbon-doped silicon semiconductor film is deposited with a chemical vapor deposition technique using the precursors silane and methane.

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Referring to FIG. 3C, a wet etch mixture **320** may be applied to the surface of semiconductor film **310**. A suitable wet etch mixture **320** may be one that selectively etches/dissolves amorphous region **314** of semiconductor film **310**, as depicted by the arrow, while having a negligible effect on epitaxial region **312**. Furthermore, a suitable wet etch mixture **320** may account for local pH differences that occur near the surface of semiconductor film **310** during the dissolving (etching) of amorphous region **314**. In accordance with an embodiment of the present invention, wet etch mixture **320** is a four-component wet etch mixture comprising an oxidizing agent, an etchant, a buffer and a diluent. In one embodiment, semiconductor film **310** is comprised of a 0-0.1% carbon-doped silicon film and four-component wet etch mixture **320** is comprised of 180 parts per volume nitric acid (70% aqueous solution), 1 part per volume hydrofluoric acid (49% aqueous solution), 1100 parts per volume acetic acid (100%, glacial) and no additional parts per volume water. In another embodiment, semiconductor film **310** is comprised of a 0.1-2% carbon-doped silicon film and four-component wet etch mixture **320** is comprised of 100 parts per volume nitric acid (70% aqueous solution), 1 part per volume hydrofluoric acid (49% aqueous solution), 200 parts per volume acetic acid (100%, glacial) and 50 additional parts per volume water. In another embodiment, semiconductor film **310** is comprised of greater than 2% carbon-doped silicon film and four-component wet etch mixture **320** is comprised of 75 parts per volume nitric acid (70% aqueous solution), 1 part per volume hydrofluoric acid (49% aqueous solution), 100 parts per volume acetic acid (100%, glacial) and 25 additional parts per volume water. In one embodiment, wet etch mixture **320** is applied in the temperature range of 20-30° C. during removal of amorphous region **314**. In another embodiment, wet etch mixture **320** is applied to the structure in FIG. 3B by an immersion technique or by a spray-on chemical injection technique. In one embodiment, amorphous region **314** is etched until completely removed by wet etch mixture **320**.

Referring to FIG. 3D, upon complete etching of amorphous region **314**, wet etch mixture **320** may be removed. In accordance with an embodiment of the present invention, only epitaxial region **312** (which is above crystalline portion **302**) of non-selectively deposited semiconductor film **310** remains above substrate **300** following treatment with wet etch mixture **320**. In one embodiment, crystalline portion **302** of substrate **300** has a larger lattice constant than epitaxial region **312**. In one embodiment, crystalline portion **302** of substrate **300** is comprised of silicon and epitaxial region **312** is comprised of carbon-doped silicon.

As an example of another embodiment of the present invention, FIGS. 4A-D illustrate the non-selective deposition and subsequent selective etching of a semiconductor film on a substrate comprising a crystalline portion and an amorphous portion, wherein a region in the crystalline portion is recessed to form an etched-out region in the crystalline portion. Referring to FIG. 4A, a region of single crystalline portion **402** of substrate **400** may be recessed to form etched-out region **406**. Etched out region **406** may be formed by first masking substrate **400** with a masking layer and then etching any exposed regions of crystalline portion **402** with a dry etch or wet etch treatment. In one embodiment, SF₆ or NF₃ gas in a plasma etcher is used to form etched-out region **406**. In accordance with one embodiment of the present invention, crystalline portion **402** is comprised of crystalline silicon or an epitaxial silicon region grown atop a distinct crystalline silicon substrate. In another embodiment, crystalline portion **402** is comprised of germanium or a III-V material such as but not limited to gallium nitride, gallium phosphide, gallium

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arsenide, indium phosphide or indium antimonide. In one embodiment, amorphous portion **404** is comprised of an oxide or a nitride material. In another embodiment, amorphous portion **404** is comprised of silicon nitride, silicon dioxide, silicon oxy-nitride or a high-k material such as hafnium oxide.

Referring to FIG. 4B, semiconductor film **410** may be deposited non-selectively above the crystalline portion **402**, the amorphous portion **404** and the etched-out region **406** of substrate **400**. In accordance with one embodiment of the present invention, a suitable non-selectively deposited semiconductor film **410** is one which forms an epitaxial region **412** above crystalline portion **402** and above etched-out region **406**, while forming an amorphous region **414** above amorphous portion **404**. Semiconductor film **410** may be deposited by any suitable technique. In one embodiment, semiconductor film **410** is deposited by chemical vapor deposition, physical vapor deposition or atomic layer deposition. In one embodiment, semiconductor film **410** is lattice-matched with crystalline portion **402**. In another embodiment, semiconductor film **410** is lattice-mismatched with and has a smaller lattice constant than crystalline portion **402**. In another embodiment, semiconductor film **410** is lattice-mismatched with and has a larger lattice constant than crystalline portion **402**. In an embodiment, semiconductor film **410** is a carbon-doped silicon film. In another embodiment, semiconductor film **410** is a carbon-doped silicon film in situ doped with phosphorus dopant atoms. In the case of a carbon-doped silicon film, the carbon concentration as a % of total film composition may vary depending on the required application. In one embodiment, the total carbon concentration of a carbon-doped silicon semiconductor film is 0-0.1%. In another embodiment, the total carbon concentration of a carbon-doped silicon semiconductor film is 0.1-2%. In another embodiment, the total carbon concentration of a carbon-doped silicon semiconductor film is greater than 2%. In one embodiment, a carbon-doped silicon semiconductor film is deposited with a chemical vapor deposition technique using the precursors silane and methane. In one embodiment, the top surface of semiconductor film **410** above etched out region **406** is raised above the top surface of substrate **400**.

Referring to FIG. 4C, a wet etch mixture **420** may be applied to the surface of semiconductor film **410**. A suitable wet etch mixture **420** may be one that selectively etches/dissolves amorphous region **414** of semiconductor film **410**, as depicted by the arrow, while having a negligible effect on epitaxial region **412**. Furthermore, a suitable wet etch mixture **420** may account for local pH differences that occur near the surface of semiconductor film **410** during the dissolving (etching) of amorphous region **414**. In accordance with an embodiment of the present invention, wet etch mixture **420** is a four-component wet etch mixture comprising an oxidizing agent, an etchant, a buffer and a diluent. In one embodiment, semiconductor film **410** is comprised of a 0-0.1% carbon-doped silicon film and four-component wet etch mixture **420** is comprised of 180 parts per volume nitric acid (70% aqueous solution), 1 part per volume hydrofluoric acid (49% aqueous solution), 1100 parts per volume acetic acid (100%, glacial) and no additional parts per volume water. In another embodiment, semiconductor film **410** is comprised of a 0.1-2% carbon-doped silicon film and four-component wet etch mixture **420** is comprised of 100 parts per volume nitric acid (70% aqueous solution), 1 part per volume hydrofluoric acid (49% aqueous solution), 200 parts per volume acetic acid (100%, glacial) and 50 additional parts per volume water. In another embodiment, semiconductor film **410** is comprised of greater than 2% carbon-doped silicon film and four-compo-

ment wet etch mixture **420** is comprised of 75 parts per volume nitric acid (70% aqueous solution), 1 part per volume hydrofluoric acid (49% aqueous solution), 100 parts per volume acetic acid (100%, glacial) and 25 additional parts per volume water. In one embodiment, wet etch mixture **420** is applied in the temperature range of 20-30° C. during removal of amorphous region **414**. In another embodiment, wet etch mixture **420** is applied to the structure in FIG. 4B by an immersion technique or by a spray-on chemical injection technique. In one embodiment, amorphous region **414** is etched until completely removed by wet etch mixture **420**.

Referring to FIG. 4D, upon complete etching of amorphous region **414**, wet etch mixture **420** may be removed. In accordance with an embodiment of the present invention, only epitaxial region **412** (which is above crystalline portion **402** and above etched-out region **406**) of non-selectively deposited semiconductor film **410** remains in and above substrate **400** following treatment with wet etch mixture **420**. In one embodiment, crystalline portion **402** of substrate **400** has a larger lattice constant than epitaxial region **412**. In another embodiment, crystalline portion **402** of substrate **400** is comprised of silicon and epitaxial region **412** is comprised of carbon-doped silicon. In one embodiment, the top surface of epitaxial region **412** in etched-out region **406** is raised above the top surface of substrate **400**.

The method to selectively etch with a four-component etch mixture, and hence pattern, a semiconductor film deposited non-selectively may be utilized in the fabrication of a semiconductor device. In one embodiment, the semiconductor device is a MOS-FET, a bipolar transistor, a memory transistor or a micro-electronic machine (MEM). In another embodiment, the semiconductor device is a planar device or a non-planar device, such as a tri-gate or double-gate transistor. For illustrative purposes, the fabrication of an NMOS-FET device incorporating a four-component selective etch process to pattern a semiconductor film deposited non-selectively is described below, in accordance with one embodiment of the present invention.

FIGS. 5A-E illustrate a process flow for forming an NMOS-FET device with strain-inducing source/drain regions as facilitated by a four-component selective etch mixture, in accordance with an embodiment of the present invention. Referring to FIG. 5A, a non-strained NMOS-FET **550** is first formed. Non-strained NMOS-FET **550** may be comprised of a channel region **506** in a crystalline substrate **500**. In one embodiment of the present invention, crystalline substrate **500** is comprised of single crystalline silicon. In another embodiment, crystalline substrate **500** is comprised of an epitaxial silicon layer grown atop a distinct crystalline silicon substrate. In one embodiment, crystalline substrate **500** is comprised of germanium or a III-V material such as but not limited to gallium nitride, gallium phosphide, gallium arsenide, indium phosphide or indium antimonide.

A gate dielectric layer **504** may be formed above channel region **506**. In one embodiment, gate dielectric layer **504** is formed by a thermal oxidation process and is comprised of silicon dioxide or silicon oxy-nitride. In another embodiment, gate dielectric layer **504** is formed by chemical vapor deposition or atomic layer deposition and is comprised of a high-k dielectric layer such as, but not limited to, hafnium oxide, zirconium oxide, hafnium silicate, hafnium oxy-nitride or lanthanum oxide.

A gate electrode **508** may be formed above gate dielectric layer **504**. Gate electrode **508** may be formed by a subtractive etching process scheme or by a replacement gate process scheme. In one embodiment, gate electrode **508** is comprised of a polycrystalline silicon gate electrode, wherein the charge

carrier dopant impurities are implanted during fabrication of the tip and source/drain regions, described below. In another embodiment, gate electrode **508** is comprised of a metal layer such as but not limited to metal nitrides, metal carbides, hafnium, zirconium, titanium, tantalum, aluminum, ruthenium, palladium, platinum, cobalt, nickel or conductive metal oxides, e.g. ruthenium oxide. In one embodiment, an amorphous gate protecting layer **540** comprised of silicon dioxide, silicon nitride, silicon oxy-nitride or carbon-doped silicon nitride is formed above gate electrode **508**, as depicted in FIG. 5A.

A tip extension **512** may be formed by implanting charge carrier dopant impurity atoms into crystalline substrate **500**. Gate electrode **508** may act to mask a portion of crystalline substrate **500** to form self-aligned tip extensions **512**. In one embodiment, boron, arsenic, phosphorus, indium or a combination thereof is implanted into crystalline substrate **500** to form tip extension **512**. In another embodiment, crystalline substrate **500** is implanted to form N-type tip extensions.

Gate dielectric layer **504** and gate electrode **508** may be isolated by amorphous gate isolation spacer **510**. Amorphous gate isolation spacer **510** may be formed adjacent the sidewalls of gate dielectric layer **504** and gate electrode **508** by any suitable technique. In an embodiment, amorphous gate isolation spacer **510** is formed from an amorphous insulating layer such as but not limited to silicon dioxide, silicon nitride, silicon oxy-nitride or carbon-doped silicon nitride that is deposited by a chemical vapor deposition process and subsequently dry etched. In another embodiment, the thickness of the amorphous insulating layer is selected to determine the final width of amorphous gate isolation spacer **510**. In one embodiment, amorphous gate isolation spacer **510** forms a hermetic seal with gate electrode **508** and the top surface of substrate **500** in order to encapsulate gate dielectric layer **504**.

A source/drain region **514** may be formed by implanting charge carrier dopant impurity atoms into crystalline substrate **500**. Thus, source/drain region **514** may be formed from the same material as channel region **506**. Therefore, the lattice mismatch between source/drain region **514** and channel region **506** may be negligible, resulting in effectively no strain induced on channel region **506**. Amorphous gate isolation spacer **510** and gate electrode **508** may act to shield a portion of crystalline substrate **500** during the implant step to form self-aligned source/drain regions **514**. In effect, the thickness of amorphous gate isolation spacer **510** may play a role in dictating the dimensions of source/drain region **514**. In one embodiment, boron, arsenic, phosphorus, indium or a combination thereof is implanted into crystalline substrate **500** to form source/drain regions **514**. In one embodiment, the source/drain implant step is carried out to add charge carrier dopant impurities to a polycrystalline gate electrode.

Referring to FIG. 5B, portions of crystalline substrate **500**, including source/drain regions **514**, may be removed to form etched-out region **516** in crystalline substrate **500**. Etched-out region **516** may be formed by any suitable technique, such as a dry etch or a wet etch process. In one embodiment, SF₆ or NF₃ gas in a plasma etcher is used to form etched-out region **516**. In another embodiment, protective layer **540** protects gate electrode **508** during the formation of etched-out region **516**. In one embodiment, etched-out region **516** is formed to a depth sufficient to remove the charge carrier dopant impurities implanted to form source/drain region **514**.

Referring to FIG. 5C, semiconductor film **530** may be deposited non-selectively on the surface of the structure formed in FIG. 5B, including above etched-out region **516** of crystalline substrate **500**. In accordance with one embodiment of the present invention, a suitable non-selectively

deposited semiconductor film **530** is one which forms an epitaxial region **532** above etched-out region **516** of crystalline substrate **500**, while forming an amorphous region **534** above amorphous gate protection layer **540** and amorphous gate isolation spacers **510**. Semiconductor film **530** may be deposited by any suitable technique. In one embodiment, semiconductor film **530** is deposited by chemical vapor deposition, physical vapor deposition or atomic layer deposition. In one embodiment, semiconductor film **530** is lattice-matched with crystalline substrate **500**. In another embodiment, semiconductor film **530** is lattice-mismatched with and has a smaller lattice constant than crystalline substrate **500**. In another embodiment, semiconductor film **530** is lattice-mismatched with and has a larger lattice constant than crystalline substrate **500**. In an embodiment, semiconductor film **530** is a carbon-doped silicon film. In another embodiment, semiconductor film **530** is a carbon-doped silicon film in situ doped with phosphorus dopant atoms. In the case of a carbon-doped silicon film, the carbon concentration as a % of total film composition may vary depending on the required application. In one embodiment, the total carbon concentration of a carbon-doped silicon semiconductor film is 0-0.1%. In another embodiment, the total carbon concentration of a carbon-doped silicon semiconductor film is 0.1-2%. In another embodiment, the total carbon concentration of a carbon-doped silicon semiconductor film is greater than 2%. In one embodiment, a carbon-doped silicon semiconductor film is deposited with a chemical vapor deposition technique using the precursors silane and methane. In one embodiment, the top surface of semiconductor film **530** above etched-out region **516** is raised above the top surface of crystalline substrate **500**.

Referring to FIG. **5D**, a wet etch mixture **520** may be applied to the surface of semiconductor film **530**. A suitable wet etch mixture **520** may be one that selectively etches/dissolves amorphous region **534** of semiconductor film **530**, as depicted by the arrow, while having a negligible effect on epitaxial region **532**. Furthermore, a suitable wet etch mixture **520** may account for local pH differences that occur near the surface of semiconductor film **530** during the dissolving (etching) of amorphous region **534**. In accordance with an embodiment of the present invention, wet etch mixture **520** is a four-component wet etch mixture comprising an oxidizing agent, an etchant, a buffer and a diluent. In one embodiment, semiconductor film **530** is comprised of a 0-0.1% carbon-doped silicon film and four-component wet etch mixture **520** is comprised of 180 parts per volume nitric acid (70% aqueous solution), 1 part per volume hydrofluoric acid (49% aqueous solution), 1100 parts per volume acetic acid (100%, glacial) and no additional parts per volume water. In another embodiment, semiconductor film **530** is comprised of a 0.1-2% carbon-doped silicon film and four-component wet etch mixture **520** is comprised of 100 parts per volume nitric acid (70% aqueous solution), 1 part per volume hydrofluoric acid (49% aqueous solution), 200 parts per volume acetic acid (100%, glacial) and 50 additional parts per volume water. In another embodiment, semiconductor film **530** is comprised of greater than 2% carbon-doped silicon film and four-component wet etch mixture **520** is comprised of 75 parts per volume nitric acid (70% aqueous solution), 1 part per volume hydrofluoric acid (49% aqueous solution), 100 parts per volume acetic acid (100%, glacial) and 25 additional parts per volume water. In one embodiment, wet etch mixture **520** is applied in the temperature range of 20-30° C. during removal of amorphous region **534**. In another embodiment, wet etch mixture **520** is applied to the structure in FIG. **5C** by an immersion technique or by a spray-on chemical injection technique. In

one embodiment, amorphous region **534** is etched until completely removed by wet etch mixture **520**.

Referring to FIG. **5E**, upon complete etching of amorphous region **534** of semiconductor film **530**, wet etch mixture **520** may be removed. In accordance with an embodiment of the present invention, only epitaxial region **532** of non-selectively deposited semiconductor film **530** remains above crystalline substrate **500** following treatment with wet etch mixture **520**. In one embodiment, crystalline substrate **500**, and hence channel region **506**, has a larger lattice constant than epitaxial region **532**. In one embodiment, crystalline substrate **500** is comprised of silicon and epitaxial region **532** is comprised of carbon-doped silicon. In one embodiment, epitaxial region **532** is comprised of carbon-doped silicon and is subsequently implanted with phosphorus dopant impurities in a concentration range of $2E19$ atoms/cm³- $2E21$ atoms/cm³.

Epitaxial region **532** may function as a strain-inducing region and thus NMOS-FET **590** in FIG. **5E** may comprise a strained source/drain region. Therefore, a uniaxial tensile strain, depicted by the arrows in FIG. **5E**, may be rendered on channel region **506** in NMOS-FET **590**, which can enhance electron mobility in the device. In one embodiment, the top surface of epitaxial region **532** is raised above the top surface of crystalline substrate **500**, where gate dielectric layer **504** resides, as depicted in FIG. **5E**. NMOS-FET **590** may subsequently be incorporated into an integrated circuit by conventional process steps, as known in the art.

The present invention is not limited to the formation of NMOS-FET devices with strain-inducing source/drain regions. In accordance with another embodiment of the present invention, a PMOS-FET comprising strain-inducing source/drain regions may be fabricated in a manner similar to that depicted in FIGS. **5A-E**. In an embodiment, a non-selectively deposited silicon/germanium film has an amorphous region and an epitaxial region. A four-component wet etch mixture may be tuned to accommodate the selective etching of the amorphous region of the silicon/germanium film while having negligible impact on the epitaxial region, wherein the four-component wet etch mixture comprises an oxidizing agent, an etchant, a buffer and a diluent.

Thus, a method to selectively etch, and hence pattern, a semiconductor film deposited non-selectively has been disclosed. In one embodiment, a carbon-doped silicon film is deposited non-selectively such that the film forms an epitaxial region where deposited on a crystalline surface and an amorphous region where deposited on an amorphous surface. A four-component wet etch mixture may be tuned to accommodate the selective etching of a variety of film compositions, wherein the four-component wet etch mixture comprises an oxidizing agent, an etchant, a buffer and a diluent. In one embodiment, the four-component wet etch mixture comprises nitric acid (oxidizing agent), hydrofluoric acid (etchant), acetic acid (buffer), and water (diluent).

What is claimed is:

1. A method of forming a semiconductor structure comprising:
 - forming a substrate comprising a crystalline portion and an amorphous portion;
 - depositing a semiconductor film non-selectively above said substrate, wherein said semiconductor film comprises an epitaxial region above said crystalline portion and an amorphous region above said amorphous portion; and
 - etching said semiconductor film with a etch mixture comprising an oxidizing agent, an etchant, a buffer and a diluent, wherein said amorphous region is removed from

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said amorphous portion of said substrate and said epitaxial region is retained above said crystalline portion of said substrate.

2. The method of claim 1 wherein said oxidizing agent comprises nitric acid, hydrogen peroxide or di-tert-butylperoxide.

3. The method of claim 1 wherein said wet etch mixture etches said amorphous region of said semiconductor film at least 20-fold faster than said epitaxial region.

4. The method of claim 1 wherein said crystalline portion of said substrate comprises silicon and said semiconductor film comprises carbon-doped silicon.

5. The method of claim 4 wherein said semiconductor film comprises 0.1-2% carbon-doped silicon and said wet etch mixture comprises 100 parts per volume nitric acid (70% aqueous solution), 1 part per volume hydrofluoric acid (49% aqueous solution), 200 parts per volume acetic acid (100%, glacial) and 50 additional parts per volume water.

6. A method of forming a semiconductor structure comprising:

forming a substrate comprising a crystalline portion and an amorphous portion;

forming an etched-out region in said crystalline portion of said substrate;

depositing a semiconductor film non-selectively above said substrate, wherein said semiconductor film comprises an epitaxial region above said crystalline portion and above said etched-out region and an amorphous region above said amorphous portion, and wherein said epitaxial region and said crystalline portion are lattice mismatched; and

etching said semiconductor film with a wet etch mixture comprising an oxidizing agent, an etchant, a buffer and a diluent, wherein said amorphous region is removed from said amorphous portion of said substrate and said epitaxial region is retained above said crystalline portion and said etched-out region of said substrate.

7. The method of claim 6 wherein said oxidizing agent comprises nitric acid, hydrogen peroxide or di-tert-butylperoxide.

8. The method of claim 6 wherein said wet etch mixture etches said amorphous region of said semiconductor film at least 20-fold faster than said epitaxial region.

9. The method of claim 6 wherein said crystalline portion of said substrate comprises silicon and said semiconductor film comprises carbon-doped silicon.

10. The method of claim 9 wherein said semiconductor film comprises 0.1-2% carbon-doped silicon and said wet etch mixture comprises 100 parts per volume nitric acid (70% aqueous solution), 1 part per volume hydrofluoric acid (49% aqueous solution), 200 parts per volume acetic acid (100%, glacial) and 50 additional parts per volume water.

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11. A method of forming a semiconductor device comprising:

forming a gate dielectric layer above a channel region in a crystalline substrate;

forming a gate electrode above said gate dielectric layer; forming an amorphous gate protecting layer above said gate electrode;

forming an amorphous gate isolation spacer adjacent the sidewalls of said gate electrode and said gate dielectric layer;

forming a source/drain region in said crystalline substrate; removing a portion of said crystalline substrate, including said source/drain region, to form an etched-out region in said crystalline substrate;

depositing a semiconductor film non-selectively above said etched-out region, above said amorphous gate protecting layer and above said amorphous gate isolation spacer, wherein said semiconductor film comprises an epitaxial region above said etched-out region, an amorphous region above said amorphous gate protecting layer and an amorphous region above said amorphous gate isolation spacer; and

etching said semiconductor film with a wet etch mixture comprising an oxidizing agent, an etchant, a buffer and a diluent, wherein said amorphous region is removed from said amorphous gate protecting layer and from said amorphous gate isolation spacer, while said epitaxial region is retained above said etched-out region of said crystalline substrate.

12. The method of claim 11 wherein said epitaxial region of said semiconductor film and said crystalline substrate are lattice mismatched.

13. The method of claim 12 wherein said crystalline substrate comprises silicon and said semiconductor film comprises carbon-doped silicon.

14. The method of claim 13 wherein said semiconductor film comprises 0.1-2% carbon-doped silicon and said wet etch mixture comprises 100 parts per volume nitric acid (70% aqueous solution), 1 part per volume hydrofluoric acid (49% aqueous solution), 200 parts per volume acetic acid (100%, glacial) and 50 additional parts per volume water.

15. The method of claim 11 wherein said oxidizing agent comprises nitric acid, hydrogen peroxide or di-tert-butylperoxide.

16. The method of claim 11 wherein said wet etch mixture etches said amorphous region above said amorphous gate protecting layer and said amorphous region above said amorphous gate isolation spacer at least 20-fold faster than said epitaxial region above said etched-out region.

17. The method of claim 16 wherein said epitaxial region and said crystalline substrate are lattice mismatched.

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